click by NT on 3/23/15

MEMORANDUM

TO:

Mr. Addison Rice

Anderson, Mulholland and Associates

DATE: March 20, 2015

FROM: R. Infante

FILE: 1503058

RE:

Data Validation

Air samples

SDG: 1503058C/1503058F/1503058H

SUMMARY

Full validation was performed on the data for several gas samples analyzed for 2-propanol and methanol

organic compounds by method Compendium Method TO-15: Determination Of Volatile Organic Compounds (VOCs) In Air Collected In Specially-Prepared Canisters And Analyzed By Gas Chromatography/Mass Spectrometry (GC/MS), January, 1999 and methane by ASTM method D-1946-modified. The samples were collected at the Bristol Myer Squib, Humacao, PR site on March 02, 2015 and submitted to Eurofins Air Toxics, Inc. of Folson, California that analyzed and reported the results under delivery groups (SDG) 1503058C/1503058F/1503058H.

The sample results were assessed according to USEPA data validation guidance documents in the following order of precedence: Compendium Method TO-15. Determination Of Volatile Organic Compounds (VOCs) In Air Collected In Specially-Prepared Canisters And Analyzed By Gas Chromatography/Mass Spectrometry (GC/MS), January, 1999; Validating Air Samples. Volatile Organic Analysis of Ambient Air in Canisters by Method TO-15, (SOP # HW-31. Revision #4. October, 2006; and the QC criteria of the ASTM method D-1946-modified. The QC criteria and data validation actions listed on the data review worksheets are from the primary guidance document, unless otherwise noted.

In general the data is valid as reported and may be used for decision making purposes. The data results are acceptable for use.

SAMPLES

The samples included in the review are listed below

| Client Sample ID | Lab. Sample ID | Collected Date | Matrix | Analysis |
|---------------------|----------------|-------------------|--------|----------|
| B6-AA-2 | 1503058C-01A | 03/02/2015 | Air | Methanol |
| B6-1IA | 1503058C-02A | 03/02/2015 | Air | Methanol |
| B6-2IA | 1503058C-03A | 03/02/2015 | Air | Methanol |
| B6-2IA Dup | 1503058C-04A | 03/02/2015 | Air | Methanol |

| Client Sample ID | Lab. Sample ID | Collected Date | Matrix | Analysis |
|---------------------|----------------|-------------------|--------|------------|
| B6-1SS | 1503058C-05A | 03/02/2015 | Air | Methanol |
| B6-2SS | 1503058C-06A | 03/02/2015 | Air | Methanol |
| B6-2SS Dup | 1503058C-07A | 03/02/2015 | Air | Methanol |
| B6-AA-2 | 1503058H-01A | 03/02/2015 | Air | 2-Propanol |
| B6-1IA | 1503058H-02A | 03/02/2015 | Air | 2-Propanol |
| B6-2IA | 1503058H-03A | 03/02/2015 | Air | 2-Propanol |
| B6-2IA Dup | 1503058H-04A | 03/02/2015 | Air | 2-Propanol |
| B6-AA-2 | 1503058F-01A | 03/02/2015 | Air | Methane |
| B6-1IA | 1503058F-02A | 03/02/2015 | Air | Methane |
| B6-2IA | 1503058F-03A | 03/02/2015 | Air | Methane |
| B6-2IA Dup | 1503058F-04A | 03/02/2015 | Air | Methane |
| B6-1SS | 1503058F-05A | 03/02/2015 | Air | Methane |
| B6-2SS | 1503058F-06A | 03/02/2015 | Air | Methane |
| B6-2SS Dup | 1503058F-07A | 03/02/2015 | Air | Methane |

REVIEW ELEMENTS

Sample data were reviewed for the following parameters, where applicable to the method

- o Agreement of analysis conducted with chain of custody (COC) form
- o Holding time and sample preservation
- o Gas chromatography/mass spectrometry (GC/MS) tunes
- o Initial and continuing calibrations
- Method blanks/trip blanks/field blank
- o Canister cleaning certification criteria
- Surrogate spike recovery
- o Internal standard performance and retention times
- o Field duplicate results
- Laboratory control sample/laboratory control sample duplicate (LCS/LCSD) results
- o Quantitation limits and sample results

DISCUSSION

Agreement of Analysis Conducted with COC Request

Sample reports corresponded to the analytical request designated on the chain-of-custody form.

Holding Times and Sample Preservation

Sample preservation was acceptable.

Samples analyzed within method recommended holding time.

GC/MS Tunes

The frequency and abundance of bromofluorobenzene (BFB) tunes were within the QC acceptance criteria. All samples were analyzed within the tuning criteria associated with the method.

Initial and Continuing Calibrations

VOCs - Methanol and 2-Propanol (Method TO-15)

The percent relative standard deviations (%RSDs) and response factors (RFs) of all target analytes were within the QC acceptance criteria in the initial calibration. Correlation coefficients (r²) of target analytes were within the QC acceptance criteria. Ongoing accuracy of the instrument was determined by the analysis of a continuing calibration standard.

Methane by ASTM method D-1946 (modified)

Initial and continuing calibrations meet method specific requirements. Initial calibration retention times meet method specific requirements.

Method Blank/Trip Blank/Field Blank

Target analytes were not detected in laboratory method blanks.

Summa canister met cleaning certification criteria.

No trip/field blank analyzed with this data package.

Surrogate Spike Recovery

The surrogate recoveries as per method TO-15 were within the laboratory QC acceptance limits in all samples analyzed. ASTM method for methane does not require surrogate standards.

Internal Standard Performance

VOCs - Methanol and 2-Propanol(TQ-15)

Samples were spiked with the method specified internal standard. Internal standard are performance and retention times met the QC acceptance criteria in all sample analyses and calibration standards.

Laboratory/Field Duplicate Results

Field/laboratory duplicates were analyzed as part of this data set. Target analytes meet the RPD performance criteria of +25% for analytes $5\times SQL$.

LCS/LCSD Results

VOCs and Methane

LCS/LCSD (blank spike) were analyzed by the laboratory associated with this data package. Recoveries and RPD within laboratory control limits.

Quantitation Limits and Sample Results

Dilutions were performed on TO-15 samples (see worksheet).

Calculations were spot checked.

Certification

The following samples 1503058C-01A; 1503058C-02A; 1503058C-03A; 1503058C-04A; 1503058C-05A; 1503058C-06A; 1503058C-07A; 1503058F-01A; 1503058F-02A; 1503058F-03A; 1503058F-04A; 1503058F-05A; 1503058F-06A; 1503058F-07A; 1503058H-01A; 1503058H-02A; 1503058H-03A; and 1503058H-04A were analyzed following standard procedures accepted by regulatory agencies. The quality control requirements met the methods criteria except in the occasions described in this document. The results are valid. Some of the results were qualified.

Rafael Infante

Chemist License 1888



Client Sample ID: B6-AA-2 Lab ID#: 1503058C-01A

EPA METHOD TO-15 GC/MS

| File Name: Dil. Factor: | 14030505 1.37 | Date of Collection: 3/2/15 9:38:00 Date of Analysis: 3/5/15 02:43 PM | | |
|----------------------------|----------------------|--|-----------------------|-------------------|
| Compound | Rpt. Limit (ppbv) | Amount (ppbv) | Rpt. Limit (ug/m3) | Amount (ug/m3) |
| Methanol | 68 | Not Detected | 90 | Not Detected |

| Surrogates | %Recovery | Method Limits |
|-----------------------|-----------|------------------|
| 1,2-Dichloroethane-d4 | 101 | 70-130 |
| Toluene-d8 | 100 | 70-130 |
| 4-Bromofluorobenzene | 96 | 70-130 |





Client Sample ID: B6-1IA Lab ID#: 1503058C-02A

EPA METHOD TO-15 GC/MS

| File Name: | 14030506 | Date of Collection: 3/2/15 12:05: | | |
|--------------|------------|-----------------------------------|------------|--------------|
| Dil. Factor: | 1.56 | Date of Analysis: 3/5/15 03:06 Pl | | |
| Compound | Rpt. Limit | Amount | Rpt. Limit | Amount |
| | (ppbv) | (ppbv) | (ug/m3) | (ug/m3) |
| Methanol | 78 | Not Detected | 100 | Not Detected |

| | | Method |
|-----------------------|-----------|--------|
| Surrogates | %Recovery | Limits |
| 1,2-Dichloroethane-d4 | 101 | 70-130 |
| Toluene-d8 | 99 | 70-130 |
| 4-Bromofluorobenzene | 100 | 70-130 |





Client Sample ID: B6-2IA Lab ID#: 1503058C-03A

EPA METHOD TO-15 GC/MS

| File Name: 14030507 Dil. Factor: 1.59 | | Date of Collection: 3/2/15 12:08:00 PM Date of Analysis: 3/5/15 04:08 PM | | |
|--|----------------------|---|-----------------------|----------------|
| Compound | Rpt. Limit (ppbv) | Amount (ppbv) | Rpt. Limit (ug/m3) | Amount (ug/m3) |
| Methanol | 80 | Not Detected | 100 | Not Detected |

| Surrogates | %Recovery | Method Limits |
|-----------------------|-----------|------------------|
| 1,2-Dichloroethane-d4 | 102 | 70-130 |
| Toluene-d8 | 100 | 70-130 |
| 4-Bromofluorobenzene | 101 | 70-130 |





Client Sample ID: B6-2IADup Lab ID#: 1503058C-04A

EPA METHOD TO-15 GC/MS

| File Name: Dil. Factor: | 14030508 1.63 | Date of Collection: 3/2/15 12:08 Date of Analysis: 3/5/15 04:59 P | | |
|----------------------------|----------------------|--|-----------------------|-------------------|
| Compound | Rpt. Limit (ppbv) | Amount (ppbv) | Rpt. Limit (ug/m3) | Amount (ug/m3) |
| Methanol | 82 | Not Detected | 110 | Not Detected |

| Surrogates | %Recovery | Method Limits |
|-----------------------|-----------|------------------|
| 1,2-Dichloroethane-d4 | 104 | 70-130 |
| Toluene-d8 | 100 | 70-130 |
| 4-Bromofluorobenzene | 99 | 70-130 |





Client Sample ID: B6-1SS Lab ID#: 1503058C-05A

EPA METHOD TO-15 GC/MS

| File Name: | 14030509 | Date of Collection: 3/2/15 4:35:00 | | |
|--------------|------------|------------------------------------|------------|--------------|
| Dil. Factor: | 2.35 | Date of Analysis: 3/5/15 05:27 PI | | |
| Compound | Rpt. Limit | Amount | Rpt. Limit | Amount |
| | (ppbv) | (ppbv) | (ug/m3) | (ug/m3) |
| Methanol | 120 | Not Detected | 150 | Not Detected |

| Surrogates | %Recovery | Method Limits |
|-----------------------|-----------|------------------|
| 1,2-Dichloroethane-d4 | 103 | 70-130 |
| Toluene-d8 | 98 | 70-130 |
| 4-Bromofluorobenzene | 99 | 70-130 |





Client Sample ID: B6-2SS Lab ID#: 1503058C-06A

EPA METHOD TO-15 GC/MS

| File Name: Dil. Factor: | 143300.0 | | Date of Collection: 3/2/15 3:08:00 P Date of Analysis: 3/5/15 06:20 PM | |
|----------------------------|----------------------|------------------|---|-------------------|
| Compound | Rpt. Limit (ppbv) | Amount (ppbv) | Rpt. Limit (ug/m3) | Amount (ug/m3) |
| Methanol | 110 | Not Detected | 150 | Not Detected |

| Cumanatan | W.D., | Method |
|-----------------------|-----------|--------|
| Surrogates | %Recovery | Limits |
| 1,2-Dichloroethane-d4 | 102 | 70-130 |
| Toluene-d8 | 98 | 70-130 |
| 4-Bromofluorobenzene | 101 | 70-130 |





Client Sample ID: B6-2SSDup Lab ID#: 1503058C-07A

EPA METHOD TO-15 GC/MS

| File Name: | 14030511 | | V15 3:08:00 PM | |
|--------------|------------|--------------|----------------|--------------|
| Dil. Factor: | 2.43 | | I5 06:40 PM | |
| Compound | Rpt. Limit | Amount | Rpt. Limit | Amount |
| | (ppbv) | (ppbv) | (ug/m3) | (ug/m3) |
| Methanol | 120 | Not Detected | 160 | Not Detected |

| | | Method | |
|-----------------------|-----------|--------|--|
| Surrogates | %Recovery | Limits | |
| 1,2-Dichloroethane-d4 | 103 | 70-130 | |
| Toluene-d8 | 100 | 70-130 | |
| 4-Bromofluorobenzene | 102 | 70-130 | |



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Page of I poliection, handling, or shipping of samples, D.O.T. Hotline (800) 467-4922 Project Manager Late Use Only Project Info: **Turn Around** Time: Pressurized by ☐ Normal Date: Rush Project # State LY Zip 10577 Pressurization Gas. 914-391-04008x Project Name Canister Pressure/Vacuum Date Time Let ID Field Sample I.D. (Location) Can # of Collection of Collection **Analyses Requested** Initial Final see below 208 - 2IADUD 208 1635 1508 B6-255000 30 1508 Hold For Analysia 30" Relinquished by: (signature) Date/Time Recoved by: (signature) Date/Time Analy de fer: Acetone, Benzare, 15:1100 1201 34.15 Relinguished by: (signature) Date/Time Received by: (signature) Ibenzene, Isoprepy Alcold, Date/Time Relinquished by: (signature) Date/Time Received by: (signature) Condition Custogy Seals Intact? Work Order # lemo (°C)... 773032321463 A75001 Yes No None 1503058



Client Sample ID: B6-AA-2 Lab ID#: 1503058H-01A

EPA METHOD TO-15 GC/MS FULL SCAN

| File Name: | p030907 | Date of Collection: 3/2/15 9: | | |
|--------------|------------|-------------------------------|------------|---------|
| Dil. Factor: | 1.37 | Date of Analysis: 3/9/15 12:2 | | |
| Compound | Rpt. Limit | Amount | Rpt. Limit | Amount |
| | (ppbv) | (ppbv) | (ug/m3) | (ug/m3) |
| 2-Propanol | 2.7 | 0.36 J | 6.7 | 0.90 J |

J = Estimated value.

| | | Method |
|-----------------------|-----------|--------|
| Surrogates | %Recovery | Limits |
| 1,2-Dichloroethane-d4 | 99 | 70-130 |
| Toluene-d8 | 104 | 70-130 |
| 4-Bromofluorobenzene | 105 | 70-130 |





Client Sample ID: B6-1IA Lab ID#: 1503058H-02A

EPA METHOD TO-15 GC/MS FULL SCAN

| File Name: Dil. Factor: | p030910 10.4 | Date of Collection: 3/2/15 12:05 Date of Analysis: 3/9/15 01:31 F | | |
|----------------------------|----------------------|---|-----------------------|-------------------|
| Compound | Rpt. Limit (ppbv) | Amount (ppbv) | Rpt. Limit (ug/m3) | Amount (ug/m3) |
| 2-Propanol | 21 | 1600 | 51 | 3900 |

| | | Method |
|-----------------------|-----------|--------|
| Surrogates | %Recovery | Limits |
| 1,2-Dichloroethane-d4 | 104 | 70-130 |
| Toluene-d8 | 99 | 70-130 |
| 4-Bromofluorobenzene | 101 | 70-130 |





Client Sample ID: B6-2IA Lab ID#: 1503058H-03A

EPA METHOD TO-15 GC/MS FULL SCAN

| File Name: Dil. Factor: | p030908 6.37 | Date of Collection: 3/2/15 12:08: Date of Analysis: 3/9/15 12:43 P | | |
|----------------------------|----------------------|--|-----------------------|-------------------|
| Compound | Rpt. Limit (ppbv) | Amount (ppbv) | Rpt. Limit (ug/m3) | Amount (ug/m3) |
| 2-Propanol | 13 | 810 | 31 | 2000 |

| | | Method | |
|-----------------------|-----------|--------|--|
| Surrogates | %Recovery | Limits | |
| 1,2-Dichloroethane-d4 | 97 | 70-130 | |
| Toluene-d8 | 100 | 70-130 | |
| 4-Bromofluorobenzene | 103 | 70-130 | |





Client Sample ID: B6-2IADup Lab ID#: 1503058H-04A

EPA METHOD TO-15 GC/MS FULL SCAN

| File Name: Dil. Factor: | p030909 6.53 | Date of Collection: 3/2/15 12:0 Date of Analysis: 3/9/15 01:07 | | |
|----------------------------|----------------------|---|-----------------------|-------------------|
| Compound | Rpt. Limit (ppbv) | Amount (ppbv) | Rpt. Limit (ug/m3) | Amount (ug/m3) |
| 2-Propanol | 13 | 870 | 32 | 2100 |

| | | Method |
|-----------------------|-----------|--------|
| Surrogates | %Recovery | Limits |
| 1,2-Dichloroethane-d4 | 97 | 70-130 |
| Toluene-d8 | 102 | 70-130 |
| 4-Bromofluorobenzene | 104 | 70-130 |



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Page 1 of 1

| Project Manager Terry Taylor Project Info: | | | | | | , 4022 | Turn | Around | Lab Use | Only | |
|---|---|-----------------------|---------------------|-------------------|---|----------------|--------------|----------------|---------------------|------------------|--|
| Collected I | collected by: (Print and Sign) Terry Taylor | | | • | | | Time: | | Pressurized by: | | |
| Company | AM A± Email | | | P.O. # | · · · | - | ☐ Normal | | Date: | | |
| Address 2 | 760 Westchale Autoity Purchase | State UY Zip KO | 577 | Projec | t# | | P P R | | Pressurization Gas: | | Gas: |
| | 14-391-0400ax | | | Projec | t Name | | | day specify | | N ₂ H | u (1) 10 11 11 11 11 11 11 11 11 11 11 11 11 |
| | | | D | ate | Time | | | Canis | ter Pres | sure/Vac | uum |
| Lab IID. | Field Sample I.D. (Location) | Can # | of Co | llection | of Collection | Analyses Reque | sted | Initial | Final | Receipt | Final |
| GIA | B6-AA-2 | 34188 | 3/2 | /15 | 0938 | see below | | 30" | 0" | | |
| CAA | B6-11A | 94305 | 3/2 | /15 | 1205 | | | 30" | | | |
| CBA | B6-21 A | 34011 | 3/2 | /15 | 1208 | | | 30" | 6" | | |
| CUL | B6-ZIADUP | 35132 | 3/2 | /15 | 1208 | | | 30 | 6" | | |
| | B6-155 | 36415 | 3/2 | /15 | 1635 | | | 30" | 54 | | |
| | B6-255 | 33398 | 3/2 | 15 | 1508 | | | 30" | 4" | | |
| | B6-255000 | 12379 | 3/2 | 15 | 1508 | 4 | | 30 | 6" | | |
| | B6-AA-3' | 35142 | 3/3 | , | -10000000000000000000000000000000000000 | Hold For Anal | VSIA | 30" | 5" | | |
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| | , | | | | | | | | | | |



Client Sample ID: B6-AA-2 Lab ID#: 1503058F-01A

NATURAL GAS ANALYSIS BY MODIFIED ASTM D-1946

| File Name: Dil. Factor: | 9030505 1.37 | | tion: 3/2/15 9:38:00 AM sis: 3/5/15 01:37 PM |
|----------------------------|-----------------|-------------------|---|
| Compound | | Rpt. Limit (%) | Amount (%) |
| Methane | | 0.00014 | 0.00015 |





Client Sample ID: B6-1IA Lab ID#: 1503058F-02A

NATURAL GAS ANALYSIS BY MODIFIED ASTM D-1946

| File Name: | 9030506 | Date of Collec | tion: 3/2/15 12:05:00 PM |
|--------------|---------|----------------|------------------------------|
| Dil. Factor: | 1.56 | | sis: <i>3/5</i> /15 01:59 PM |
| | | Rpt Limit | Amount |
| Compound | | (%) | (%) |
| Methane | | 0.00016 | 0.00017 |





Client Sample ID: B6-2IA Lab ID#: 1503058F-03A

NATURAL GAS ANALYSIS BY MODIFIED ASTM D-1946

| File Name: Dil. Factor: | 9030507 1.59 | | tion: 3/2/15 12:08:00 PM sis: 3/5/15 02:39 PM |
|----------------------------|-----------------|-------------------|--|
| Compound | | Rpt. Limit (%) | Amount (%) |
| Methane | | 0.00016 | 0.00018 |





Client Sample ID: B6-2IADup Lab ID#: 1503058F-04A

NATURAL GAS ANALYSIS BY MODIFIED ASTM D-1946

| File Name: Dil. Factor: | 9030508 1.63 | | tion: 3/2/15 12:08:00 PM sis: 3/5/15 04:00 PM |
|----------------------------|-----------------|------------|--|
| _ | | Rpt. Limit | Amount |
| Compound | | (%) | (%) |
| Methane | | 0.00016 | 0.00025 |





Client Sample ID: B6-1SS Lab ID#: 1503058F-05A

NATURAL GAS ANALYSIS BY MODIFIED ASTM D-1946

| File Name: | 9030509 | Date of Collec | tion: 3/2/15 4:35:00 PM |
|--------------|---------|----------------|-------------------------|
| Dil. Factor: | 2.35 | Date of Analys | sis: 3/5/15 04:33 PM |
| | | Rpt. Limit | Amount |
| Compound | | (%) | (%) |
| Methane | | 0.00024 | 0.00014 J |

J = Estimated value.





Client Sample ID: B6-2SS Lab ID#: 1503058F-06A

NATURAL GAS ANALYSIS BY MODIFIED ASTM D-1946

| File Name: Dil. Factor: | 9030510 2.24 | | tion: 3/2/15 3:08:00 PM sis: 3/5/15 05:01 PM |
|----------------------------|-----------------|-------------------|---|
| Compound | | Rpt. Limit (%) | Amount (%) |
| Methane | | 0.00022 | 0.00017 J |

J = Estimated value.



Client Sample ID: B6-2SSDup Lab ID#: 1503058F-07A

NATURAL GAS ANALYSIS BY MODIFIED ASTM D-1946

File Name: 9030511 Date of Collection: 3/2/15 3:08:00 PM
Dil. Factor: 2.43 Date of Analysis: 3/5/15 05:52 PM

Rpt. Limit Amount
Compound (%) (%)

Methane 0.00024 0.00016 J

J = Estimated value.



| | eurofins | I | Air Toxics |
|--|----------|---|------------|
|--|----------|---|------------|

| Project Manager lerry Toylor | | Project Info: | | | Turn Around | | ee Crig | |
|--|--|-----------------|----------|--------------------|---|-----------|-------------|--|
| Collected by: (Print and Sign) Terry Tentor | | | • | | | Time: | Pres | aurized by: |
| Company AM A+ | | 4 | P.O. #_ | * | *************************************** | ☐ Norma | *304.00 | n istija s |
| Address 2760 West chale Autity Richese State | OLY ZIP LOS | 233 | Project | t# | * | Rush | | surization Gas: |
| Phone 914-3-1201-04008x | | T T | Project | t Name | * | 3da | | N, He |
| Lab ED. Field Sample I.D. (Location) | Can# | Dat of Colle | | Time of Collection | Analyses Reques | | anister Pre | essure/Vacuum |
| 86-AA-2 | 34188 | 3/2/ | | 0938 | see below | 30 | | |
| B6-11A | 94305 | | 4 | 1205 | | 30 | 2" 51 | |
| B6-21A | | 3/2/ | | 1208 | | 3(| | |
| BG-2IADup | 35132 | 3/2/ | 15 | 1208 | | 30 | 5 6 | |
| B6-155 | 36415 | 3/2/ | 15 | 1635 | | 3, | 0 54 | |
| 16-255 | 33398 : | 3/2/ | 15 | 1508 | | 30 | - | 41 |
| B6-255000 | 12379 | 3/2/ | 15 | 1508 | _ V | 3 | 0 6" | |
| 86-AA-3' | 35142 | 3/3/ | 15 | 0945 | Hold For Analy | sia 30 | 5" 5" | |
| | | | | | 1 | | | |
| | 10 | | j | | , | | | |
| 3/3/15:1100 | Med bly: (signatur | 82 3 | ate/Time | | Notes: Analy 2 | Ev: A | cetone | e, Benzene, |
| 1.0 | ived by: (signatur | re) Da | ate/Time | 9 | EU, Iber | a seve, l | suprep. | y Alcohol, luene, Report to MOL |
| | ived by: (signatur | re) Dat | ite/Time | 3 | Yylene, an | e Mell | hend. | Report to |
| Leto Shipped Hearter Air Rail # | THE RESERVE OF THE PARTY OF THE | mp (°C) | | Condition. | | | | |
| Only Fed Ex 7730323214 | 163 N | A | | Errod | Yes / No | (None | J 150 | 03058 |
| | * | | | , | | | | |

| | Project Number:1503058C |
|--|--|
| | Date:02/03/2015 |
| REVIEW OF VOLATILE ORGA The following guidelines for evaluating volatile organics w | vere created to delineate required validation |
| actions. This document will assist the reviewer in using producision and in better serving the needs of the data users. The | rofessional judgment to make more informed he sample results were assessed according to |
| USEPA data validation guidance documents in the follow "Compendium Method TO-15. Determination of Volatile Organical Compension of V | wing order of precedence: QC chiefla from ganic Compounds (VOCs) In Air Collected In |
| Specially-Prepared Canisters and Analyzed By Gas Ch | nromatography/Mass Spectrometry (GC/MS), |
| January, 1999, USEPA Hazardous Waste Support Brand Analysis of Ambient Air in Canisters by Method TO-15, (SOI | ch. Validating Air Samples. Volatile Organic D# HW.31 Revision #4 October 2006) The |
| QC criteria and data validation actions listed on the data rev | iew worksheets are from the primary guidance |
| document, unless otherwise noted. | |
| The hardcopied (laboratory name) _EurofinsAir_Toxics reviewed and the quality control and performance data summ | data package received has been |
| reviewed and the quality control and performance data summ | ialized. The data feview for voos included. |
| Lab. Project/SDG No.:1503058C | Sample matrix:Air |
| No. of Samples:7 | |
| Trip blank No.: | |
| Field blank No.: | |
| Equipment blank No.: | |
| Field duplicate No.:B6-2IA/B6-2IA_Dup;_B6-2SS/B6 | -2SS_Dup |
| X Data Completeness | X Laboratory Control Spikes |
| X Holding Times | X Field Duplicates |
| X GC/MS Tuning | X Calibrations |
| X Internal Standard Performance | X Compound Identifications |
| XBlanks | X Compound Quantitation X Quantitation Limits |
| X Surrogate Recoveries N/A_ Matrix Spike/Matrix Spike Duplicate | A Quantitation Limits |
| | |
| Overall Comments:_Methanol_by_method_TO-15 | |
| Definition of Qualifiers: | |
| J- Estimated results | |
| U- Compound not detected R- Rejected data | |
| UJ- Estimated nondetect | |
| Roll Mit | |
| Reviewer: / \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ | |
| Date VJ/2V/2V17 | |

DATA COMPLETENESS

| MISSING INFORMATION | DATE LAB. CONTACTED | DATE RECEIVED |
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| All criteria were metX |
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| Criteria were not met |
| and/or see below |

HOLDING TIMES

The objective of this parameter is to ascertain the validity of the results based on the holding time of the sample from time of collection to the time of analysis.

Complete table for all samples and note the analysis and/or preservation not within criteria

| SAMPLE ID | DATE SAMPLED | DATE ANALYZED | pН | ACTION |
|-----------|------------------------|-----------------------|----------|--------------|
| | | | | |
| , | All samples analyzed w | ithin the recommended | l method | holding time |
| | | | | |
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Criteria

Aqueous samples – 14 days from sample collection for preserved samples (pH \leq 2, 4°C), no air bubbles.

Aqueous samples – 7 days from sample collection for unpreserved samples, 4°C, no air bubbles. Soil samples- 7 days from sample collection.

Cooler temperature (Criteria: 4 ± 2 °C): N/A – summa canisters

Actions

If the VOCs vial(s) have air bubbles, estimate positive results (J) and reject nondetects (R).

If the % solids of soil samples is 10-50%, estimates positive results (J) and nondetects (UJ)

If the % solid of soil samples is < 10%, estimate positive results (J) and reject nondetects (R).

If holding times are exceeded but < 14 days beyond criteria, estimate positive results (J) and nondetects (UJ).

If holding times are exceeded but < 28 days beyond criteria, estimate positive results (J) and reject nondetects (R).

If holding times are grossly exceeded (> 28 days beyond criteria), reject all results (R).

If samples were not iced or if the ice were melted (> 10°C), estimate positive results (J) and nondetects (UJ).

| | | 0- | All criteria were metX |
|--|-------------------------|-----------------------------------|-------------------------------|
| | | Cr | iteria were not met see below |
| GC/MS TUNING | | | |
| The assessment of standard tuning QC | | o determine if the sample instru | umentation is within the |
| XThe BFB p | erformance results were | e reviewed and found to be within | the specified criteria. |
| XBFB tuning | was performed for ever | ry 24 hours of sample analysis. | |
| lf no, use professi qualified or rejected | | nine whether the associated da | ta should be accepted, |
| List | the | samples | affected: |
| | | 4 | |

If mass calibration is in error, all associated data are rejected.

4

| All criteria were metX |
|------------------------|
| Criteria were not met |
| and/or see below |

CALIBRATION VERIFICATION

Compliance requirements for satisfactory instrument calibration are established to ensure that the instrument is capable of producing and maintaining acceptable quantitative data.

| Date of initial calibration: | _03/05/15 |
|----------------------------------|-----------|
| Dates of continuing calibration: | 03/05/15 |
| Instrument ID numbers:MS | SD-14 |
| Matrix/Level:/ | \ir/low |

| DATE | LAB | FILE | CRITERIA OUT | COMPOUND | SAMPLES |
|------|-----|------|--|---|---|
| | ID# | | RFs, %RSD, %D, r | | AFFECTED |
| | | | | | |
| | | | Initial and continuing comes meet method specification | alibrations meet method sic requirements. | pecific requirements. |
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Criteria

All RFs must be > 0.05 regardless of method requirements for SPCC.

All %RSD must be \leq 15 % regardless of method requirements for CCC.

All %Ds must be < 30% regardless of method requirements for CCC.

Method TO-15 does not specify criterion for the curve correlation coefficient (r). A limit for r of \geq 0.995 has therefore been utilized as professional judgment.

Actions

If any compound has an initial RF or a continuing RF of < 0.05, estimate positive results (J) and reject nondetects (R), regardless of method requirements.

If any compound has a %RSD > 15%, estimate positive results (J) and use professional judgment to qualify nondetects.

If any compound has a %RSD > 90%, estimate positive results (J) and reject nondetects (R).

If any compound has a % D > 30%, estimate positive results (J) and reject nondetects (R).

If any compound has a % D > 30%, estimate positive results (J) and nondetects (UJ).

If any compound has a % D > 90%, estimate positive results (J) and reject nondetects (R).

If any compound has r < 0.995, estimate positive results and nondetects.

A separate worksheet should be filled for each initial curve

| All criteria were metX |
|------------------------|
| Criteria were not met |
| and/or see below |

V A. BLANK ANALYSIS RESULTS (Sections 1 & 2)

The assessment of the blank analysis results is to determine the existence and magnitude of contamination problems. The criteria for evaluation of blanks apply only to blanks associated with the samples, including trip, equipment, and laboratory blanks. If problems with any blanks exist, all data associated with the case must be carefully evaluated to determine whether or not there is an inherent variability in the data for the case, or if the problem is an isolated occurrence not affecting other data.

List the contamination in the blanks below. High and low levels blanks must be treated separately.

Laboratory blanks

| DATE ANALYZED | LAB ID | LEVEL/ MATRIX | COMPOUND | CONCENTRATION UNITS |
|-------------------|-----------------|--------------------|--------------------|---------------------|
| All_method | d_blank_meeth | _method_specif | fic_criteria | |
| Summa_ca | anisters_met_cl | eaning_certifica | | |
| Field/Equipmen | t/Trip blank | | | |
| DATE ANALYZED | LAB ID | LEVEL/ Matrix | COMPOUND | CONCENTRATION UNITS |
| No_field/trip/equ | uipment_blanks | _analyzed_with | _this_data_package | |
| | | | | |
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| All criteria were metX |
|------------------------|
| Criteria were not met |
| and/or see below |

VB. BLANK ANALYSIS RESULTS (Section 3)

Blank Actions

Action Levels (ALs) should be based upon the highest concentration of contaminant determined in any blank. Do not qualify any blank with another blank. The ALs for samples which have been diluted should be corrected for the sample dilution factor and/or % moisture, where applicable. No positive sample results should be reported unless the concentration of the compound in the samples exceeds the ALs:

ALs = 10x the amount of common contaminants (methylene chloride, acetone, 2-butanone, and toluene)

ALs = 5x for any other compounds

Specific actions are as follows:

If the concentration is < sample quantitation limit (SQL) and \le AL, report the compound as not detected (U) at the SQL.

If the concentration is \geq SQL but \leq AL, report the compound as not detected (U) at the reported concentration.

If the concentration is \geq SQL and > AL, report the concentration unqualified.

Notes:

High and low level blanks must be treated separately

Compounds qualified "U" for blank contamination are still considered "hits" when qualifying for calibration criteria.

| CONTAMINATION SOURCE/LEVEL | COMPOUND | CONC/UNITS | AL/UNITS | SQL | AFFECTED SAMPLES |
|----------------------------|----------|---|----------|-----|---------------------|
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| All criteria were metX | |
|------------------------|--|
| Criteria were not met | |
| and/or see below | |

SURROGATE SPIKE RECOVERIES

Laboratory performance of individual samples is established by evaluation of surrogate spike recoveries. All samples are spiked with surrogate compounds prior to sample analysis. The accuracy of the analysis is measured by the surrogate percent recovery. Since the effects of the sample matrix are frequently outside the control of the laboratory and may present relatively unique problems, the validation of data is frequently subjective and demands analytical experience and professional judgment.

List the percent recoveries (%Rs) which do not meet the criteria for surrogate recovery. Matrix: solid/aqueous

SAMPLE ID

SURROGATE COMPOUND

ACTION

1,2-DICHLOROETHANEd4

Toluene- 4-BFB

d8

| _Surrogate_recoveries_within_laboratory_control_limits | | | | | |
|--|-------------------|--|--|--|--|
| | | | | | |
| | | | | | |
| | | | | | |
| QC Limits* (Air) | | | | | |
| LL_to_UL70to_130 | _70to_13070to_130 | | | | |

- * QC limits are laboratory in-house performance criteria, LL = lower limit, UL = upper limit.
- * If QC limits are not available, use limits of 80 120 % for aqueous and 70 130 % for solid samples.

Actions:

| QUALITY | %R < 10% | %R = 10% - LL | %R > UL |
|--------------------|----------|---------------|---------|
| Positive results | J | J | J |
| Nondetects results | R | UJ | Accept |

Surrogate action should be applied:

If one or more surrogate in the VOC fraction is out of specification, but has a recovery of > 10%.

If any one surrogate in a fraction shows < 10 % recovery.

| All criteria were met |
|-----------------------|
| Criteria were not met |
| and/or see belowN/A |

VII. A MATRIX SPIKE/MATRIX SPIKE DUPLICATE (MS/MSD)

This data is generated to determine long term precision and accuracy in the analytical method for various matrices. This data alone cannot be used to evaluate the precision and accuracy of individual samples. If any % R in the MS or MSD falls outside the designated range, the reviewer should determine if there are matrix effects, i.e. LCS data are within the QC limits but MS/MSD data are outside QC limit.

1. MS/MSD Recoveries and Precision Criteria

The laboratory should use one MS and a duplicate analysis of an unspiked field sample if target analytes are expected in the sample. If target analytes are not expected, MS/MSD should be analyzed.

| List the %Rs, RPD of the compounds which d Sample ID: | | | o not meet the criteria. Matrix/Level: | | | |
|--|------------------------|-----|---|-----------|----------------------------|--|
| MS OR MSD | COMPOUND | % R | RPD | QC LIMITS | ACTION | |
| | are_not_required_as | | | | ike_used_to_assess | |
| | s are laboratory in-ho | • | | • | r limit, UL = upper limit. | |

Actions:

| QUALITY | %R < LL | %R > UL |
|--------------------|---------|---------|
| Positive results | J | J |
| Nondetects results | R | Accept |

MS/MSD criteria apply only to the unspiked sample, its dilutions, and the associated MS/MSD samples:

If the % R for the affected compounds were < LL (or 70 %), qualify positive results (J) and nondetects (UJ).

If the % R for the affected compounds were > UL (or 130 %), only qualify positive results (J).

If 25 % or more of all MS/MSD %R were < LL (or 70 %) or if two or more MS/MSD %Rs were < 10%, qualify all positive results (J) and reject nondetects (R).

A separate worksheet should be used for each MS/MSD pair.

| All criteria were met |
|-----------------------|
| Criteria were not met |
| and/or see below N/A_ |

VII. B MATRIX SPIKE/MATRIX SPIKE DUPLICATE

MS/MSD - Unspiked Compounds

It should be noted that Method TO-15 does not specify a MS/MSD criteria for the unspiked compounds in the sample. A %RSD of < 50% has therefore been utilized as professional judgment.

If all target analytes were spiked in the MS/MSD, this review element is not applicable.

List the %RSD of the compounds which do not meet the criteria.

| Sample ID: | | | Matrix/Level/Unit: | | |
|------------|--|----------|--------------------|--|--------------|
| COMPOUND | SAMPLE CONC. | MS CONC. | MSD CONC. | % RSD | ACTION |
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Actions:

^{*} If the % RSD > 50, qualify the positive result in the unspiked samples as estimated (J).

^{*} If the % RSD is not calculated (NC) due to nondetected value, use professional judgment to qualify the data.

| All criteria were met |
|-----------------------|
| Criteria were not met |
| and/or see below N/A |

VIII. LABORATORY CONTROL SAMPLE (LCS) ANALYSIS

This data is generated to determine accuracy of the analytical method for various matrices.

1. LCS Recoveries Criteria

Where LCS spiked with the same analyte at the same concentrations as the MS/MSD? Yes or No. If no make note in data review memo.

List the %R of compounds which do not meet the criteria

| | LCS ID | COMPOUND | % R | QC LIMIT |
|----------|----------------|----------|-----|---------------------------------------|
| LCS/LCSE | D_not_required | | | · · · · · · · · · · · · · · · · · · · |
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- * QC limits are laboratory in-house performance criteria, LL = lower limit, UL = upper limit.
- * If QC limits are not available, use limits of 70 130 %.

Actions:

| QUALITY | %R < LL | %R > UL |
|--------------------|---------|---------|
| Positive results | J | J |
| Nondetects results | R | Accept |

All analytes in the associated sample results are qualified for the following criteria.

If 25 % of the LCS recoveries were < LL (or 70 %), qualify all positive results (j) and reject nondetects (R).

If two or more LCS were below 10 %, qualify all positive results as (J) and reject nondetects (R).

2. Frequency Criteria:

Where LCS analyzed at the required frequency and for each matrix? <u>Yes</u> or No. If no, the data may be affected. Use professional judgment to determine the severity of the effect and qualify data accordingly. Discuss any actions below and list the samples affected.

| | | | All criteria were met Criteria were not met and/or see belowN/A |
|-----|----------------------------|--|---|
| IX. | LABORATOR | Y DUPLICATE PRECISION | |
| | Sample IDs: Sample IDs: | _B6-2IA/B6-2IA_Dup _B6-2SS/B6-2SS_Dup | Matrix:Air Matrix:Air |

Field duplicates samples may be taken and analyzed as an indication of overall precision. These analyses measure both field and lab precision; therefore, the results may have more variability than laboratory duplicates which only laboratory performance. It is also expected that soil duplicate results will have a greater variance than water matrices due to difficulties associated with collecting identical field duplicate samples.

The project QAPP should be reviewed for project-specific information. Suggested criteria: RPD \pm 25% for air samples. If both samples and duplicate are <5 SQL, the RPD criteria is doubled.

| COMPOUND | SQL | SAMPLE CONC. | DUPLICATE CONC. | RPD | ACTION |
|----------|-------|----------------|-----------------|------------|--------|
| 1 | | | | | |
| | RPD v | vithin the met | hod performand | e criteria | |
| | | | | | |
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Actions:

Qualify as estimated positive results (J) and nondetects (UJ) for the compound that exceeded the above criteria. For organics, only the sample and duplicate will be qualified.

If an RPD cannot be calculated because one or both of the sample results is not detected, the following actions apply:

If one sample result is not detected and the other is greater than 5x the SQL qualify (J/UJ).

If one sample value is not detected and the other is greater than 5x the SQL and the SQLs for the sample and duplicate are significantly different, use professional judgment to determine if qualification is appropriate.

If one sample value is not detected and the other is less than 5x, use professional judgment to determine if qualification is appropriate.

If both sample and duplicate results are not detected, no action is needed.

| All criteria were metX |
|------------------------|
| Criteria were not met |
| and/or see below |

X. INTERNAL STANDARD PERFORMANCE

The assessment of the internal standard (IS) parameter is used to assist the data reviewer in determining the condition of the analytical instrumentation.

List the internal standard area of samples which do not meet the criteria.

- * Area of +40% or -40% of the IS area in the associated calibration standard.
- * Retention time (RT) within \pm 0.06 seconds of the IS area in the associated calibration standard.

| DATE | SAMPLE ID | IS OUT | IS AREA | ACCEPTABLE RANGE | ACTION |
|------|-------------------------------------|--------|---------------------------------------|----------------------|--------------|
| | tandard_area_and_reration_standards | | | _control_limits_for_ | both_samples |
| | | | | | |
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Actions:

1. IS actions should be applied to the compound quantitated with the out-of-control ISs

| QUALITY | IS AREA < -40% | IS AREA > + 40% |
|---------------------|----------------|-----------------|
| Positive results | J | J |
| Nondetected results | R | ACCEPT |

2. If a IS retention time varies more than 0.330 seconds, the chromatographic profile for that sample must be examined to determine if any false positive or negative exists. For shifts of a large magnitude, the reviewer may consider partial or total rejection of the data for the sample fraction.

| All criteria were metX |
|------------------------|
| Criteria were not met |
| and/or see below |

XII. SAMPLE QUANTITATION

The sample quantitation evaluation is to verify laboratory quantitation results. In the space below, please show a minimum of one sample calculation:

Calibration check 03/05/15

Methanol

RF = 25.10039

[] = (243204)(400)/(77514)(25.10039)

= 50.0 ppbv OK

| All criteria were met _X_ |
|---------------------------|
| Criteria were not met |
| and/or see below |

XII. QUANTITATION LIMITS

A. Dilution performed

| SAMPLE ID | DILUTION FACTOR | REASONS FOR DILUTION |
|----------------|----------------------------|----------------------|
| Samples dilute | d by a factor of less than | 2.50 |
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| Percent Solids | |
|---------------------------------------|-----|
| List samples which have ≤ 50 % solids | |
| | 200 |
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| | 333 |

Actions:

If the % solids of a soil sample is 10-50%, estimate positive results (J) and nondetects (UJ)

If the % solids of a soil sample is < 10%, estimate positive results (J) and reject nondetects (R)

| | Project Number:1503058H |
|--|---|
| | Date:02/03/2015 |
| | 1110 D10K10E |
| REVIEW OF VOLATILE ORG | |
| The following guidelines for evaluating volatile organics v | were created to delineate required validation |
| actions. This document will assist the reviewer in using p | to expelle regulte were assessed according to |
| lecision and in better serving the needs of the data users. T JSEPA data validation guidance documents in the follo | tie sample results were assessed according to |
| OsePA data validation guidance documents in the folioon Compendium Method TO-15. Determination of Volatile Or | apple Compounds (VOCs) In Air Collected in |
| Specially-Prepared Canisters and Analyzed By Gas Cl | promatography/Mass Spectrometry (GC/MS). |
| lanuary, 1999"; USEPA Hazardous Waste Support Bran | ch Validating Air Samples Volatile Organic |
| Analysis of Ambient Air in Canisters by Method TO-15, (SO | P # HW-31, Revision #4, October, 2006). The |
| QC criteria and data validation actions listed on the data rev | riew worksheets are from the primary guidance |
| document, unless otherwise noted. | , , , |
| The hardcopied (laboratory name) _EurofinsAir_Toxics | data package received has been |
| eviewed and the quality control and performance data summ | narized. The data review for VOCs included: |
| • • • • | |
| _ab. Project/SDG No.:1503058H | Sample matrix:Air |
| No. of Samples:4 | |
| | |
| Trip blank No.: | |
| Field blank No.: | |
| =quipment diank ivo.: | |
| Field duplicate No.:B6-2IA/B6-2IA_Dup | |
| V Deta Osmalatanasa | V Laboratory Control Chikon |
| X Data Completeness | X Laboratory Control Spikes |
| X Holding Times | X Field Duplicates |
| X GC/MS Tuning | X Calibrations |
| X Internal Standard Performance | X Compound Identifications |
| X Blanks | X Compound Quantitation |
| X Surrogate Recoveries | X Quantitation Limits |
| N/A_ Matrix Spike/Matrix Spike Duplicate | |
| 0 110 1 0 D in a lib and TO 45 | |
| Overall Comments:_2-Propanol_by_method_TO-15 | |
| | |
| Definition of Qualifiers: | |
| J- Estimated results | |
| U- Compound not detected | |
| R- Rejected data | |
| UJ- Estimated nondetect | |
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| Kall N. A. | |
| Reviewer: / Wax Manx | - |
| Date: 03/20/2015 | |

DATA COMPLETENESS

| MISSING INFORMATION | DATE LAB. CONTACTED | DATE RECEIVED |
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| All criteria were metX |
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| Criteria were not met |
| and/or see below |

HOLDING TIMES

The objective of this parameter is to ascertain the validity of the results based on the holding time of the sample from time of collection to the time of analysis.

Complete table for all samples and note the analysis and/or preservation not within criteria

| SAMPLE ID | DATE SAMPLED | DATE ANALYZED | pН | ACTION |
|-----------|-----------------------|----------------------------|----------|--------------|
| | | | | |
| Δ | Il samples analyzed w | vithin the recommended | 1 method | holding time |
| | in Samples analyzed H | | 11160100 | nowing unic |
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<u>Criteria</u>

Aqueous samples – 14 days from sample collection for preserved samples (pH \leq 2, 4°C), no air bubbles.

Aqueous samples -7 days from sample collection for unpreserved samples, 4° C, no air bubbles. Soil samples -7 days from sample collection.

Cooler temperature (Criteria: 4 + 2 °C): N/A – summa canisters

Actions

chica i se la constitución

If the VOCs vial(s) have air bubbles, estimate positive results (J) and reject nondetects (R).

If the % solids of soil samples is 10-50%, estimates positive results (J) and nondetects (UJ)

If the % solid of soil samples is < 10%, estimate positive results (J) and reject nondetects (R).

If holding times are exceeded but < 14 days beyond criteria, estimate positive results (J) and nondetects (UJ).

If holding times are exceeded but < 28 days beyond criteria, estimate positive results (J) and reject nondetects (R).

If holding times are grossly exceeded (> 28 days beyond criteria), reject all results (R).

If samples were not iced or if the ice were melted (> 10°C), estimate positive results (J) and nondetects (UJ).

| | | (| All criteria were metX Criteria were not met see below |
|---|---------------------|-----------------------------------|---|
| GC/MS TUNING | | | |
| The assessment of th standard tuning QC lim | _ | s to determine if the sample inst | rumentation is within the |
| _X The BFB perfo | ormance results we | ere reviewed and found to be with | in the specified criteria. |
| XBFB tuning wa | as performed for ev | very 24 hours of sample analysis. | |
| If no, use professiona qualified or rejected. | l judgment to det | ermine whether the associated d | ata should be accepted, |
| List | the | samples | affected: |
| If mass calibration is in | error, all associat | ed data are rejected. | _ |

| All criteria were metX |
|------------------------|
| Criteria were not met |
| and/or see below |

CALIBRATION VERIFICATION

Compliance requirements for satisfactory instrument calibration are established to ensure that the instrument is capable of producing and maintaining acceptable quantitative data.

| Date of initial calibration: | _02/17/15 |
|----------------------------------|-----------|
| Dates of continuing calibration: | 03/09/15 |
| Instrument ID numbers:MS | SD-P |
| Matrix/Level:A | vir/low |

| DATE | LAB | FILE | CRITERIA OUT | COMPOUND | SAMPLES | | | | |
|-----------|--|----------|------------------|----------|----------|--|--|--|--|
| | ID# | | RFs, %RSD, %D, r | | AFFECTED | | | | |
| | | | | | | | | | |
| | Initial and continuing calibrations meet method specific requirements. Initial calibration retention | | | | | | | | |
| times mee | t method | specific | requirements. | | | | | | |
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Criteria

All RFs must be > 0.05 regardless of method requirements for SPCC.

All %RSD must be \leq 15 % regardless of method requirements for CCC.

All %Ds must be $\leq 30\%$ regardless of method requirements for CCC.

Method TO-15 does not specify criterion for the curve correlation coefficient (r). A limit for r of \geq 0.995 has therefore been utilized as professional judgment.

Actions

If any compound has an initial RF or a continuing RF of < 0.05, estimate positive results (J) and reject nondetects (R), regardless of method requirements.

If any compound has a %RSD > 15%, estimate positive results (J) and use professional judgment to qualify nondetects.

If any compound has a %RSD > 90%, estimate positive results (J) and reject nondetects (R).

If any compound has a % D > 30%, estimate positive results (J) and reject nondetects (R).

If any compound has a % D > 30%, estimate positive results (J) and nondetects (UJ).

If any compound has a % D > 90%, estimate positive results (J) and reject nondetects (R).

If any compound has r < 0.995, estimate positive results and nondetects.

A separate worksheet should be filled for each initial curve

| All criteria were metX |
|------------------------|
| Criteria were not met |
| and/or see below |

V A. BLANK ANALYSIS RESULTS (Sections 1 & 2)

The assessment of the blank analysis results is to determine the existence and magnitude of contamination problems. The criteria for evaluation of blanks apply only to blanks associated with the samples, including trip, equipment, and laboratory blanks. If problems with any blanks exist, all data associated with the case must be carefully evaluated to determine whether or not there is an inherent variability in the data for the case, or if the problem is an isolated occurrence not affecting other data.

List the contamination in the blanks below. High and low levels blanks must be treated separately.

Laboratory blanks

| DATE ANALYZED | LAB ID | LEVEL/ MATRIX | COMPOUND | CONCENTRATION UNITS |
|-------------------|----------------|-------------------|---------------------|---------------------|
| All_method | d_blank_meeth | _method_speci | fic_criteria | |
| Summa_ca | - | | | |
| Field/Equipmen | | And different | | |
| DATE ANALYZED | LAB ID | LEVEL/ MATRIX | COMPOUND | CONCENTRATION UNITS |
| No_field/trip/equ | uipment_blanks | _analyzed_with | n_this_data_package | |
| | | | | |
| | | | | |
| | | | | |
| | | | | |

| All criteria were metX |
|------------------------|
| Criteria were not met |
| and/or see below |

VB. BLANK ANALYSIS RESULTS (Section 3)

Blank Actions

Action Levels (ALs) should be based upon the highest concentration of contaminant determined in any blank. Do not qualify any blank with another blank. The ALs for samples which have been diluted should be corrected for the sample dilution factor and/or % moisture, where applicable. No positive sample results should be reported unless the concentration of the compound in the samples exceeds the ALs:

ALs = 10x the amount of common contaminants (methylene chloride, acetone, 2-butanone, and toluene)

ALs = 5x for any other compounds

Specific actions are as follows:

If the concentration is < sample quantitation limit (SQL) and \le AL, report the compound as not detected (U) at the SQL.

If the concentration is \geq SQL but \leq AL, report the compound as not detected (U) at the reported concentration.

If the concentration is \geq SQL and > AL, report the concentration unqualified.

Notes:

High and low level blanks must be treated separately Compounds qualified "U" for blank contamination are still considered "hits" when qualifying for calibration criteria.

| CONTAMINATION SOURCE/LEVEL | COMPOUND | CONC/UNITS | AL/UNITS | SQL | AFFECTED SAMPLES |
|----------------------------|--|------------|----------|-----|---------------------|
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| All criteria were met _X |
|--------------------------|
| Criteria were not met |
| and/or see below |

SURROGATE SPIKE RECOVERIES

Laboratory performance of individual samples is established by evaluation of surrogate spike recoveries. All samples are spiked with surrogate compounds prior to sample analysis. The accuracy of the analysis is measured by the surrogate percent recovery. Since the effects of the sample matrix are frequently outside the control of the laboratory and may present relatively unique problems, the validation of data is frequently subjective and demands analytical experience and professional judgment.

List the percent recoveries (%Rs) which do not meet the criteria for surrogate recovery. Matrix: solid/aqueous

| SAMPLE ID | SURRO | SURROGATE COMPOUND | | | | |
|------------------|-------------------------------|--------------------|------------|--|--|--|
| | 1,2-DICHLOROETHANE- d4 | Toluene- d8 | 4-BFB | | | |
| _Surrogate_rec | overies_within_laboratory_con | ntrol_limits | | | | |
| | | | | | | |
| | | | | | | |
| | | | | | | |
| QC Limits* (Air) | | | | | | |
| LL_to_U | IL70to_130 | _70to_13 | 3070to_130 | | | |

- * QC limits are laboratory in-house performance criteria, LL = lower limit, UL = upper limit.
- * If QC limits are not available, use limits of 80 120 % for aqueous and 70 130 % for solid samples.

Actions:

| QUALITY | %R < 10% | %R = 10% - LL | %R > UL |
|--------------------|----------|---------------|---------|
| Positive results | J | J | J |
| Nondetects results | R | UJ | Accept |

Surrogate action should be applied:

If one or more surrogate in the VOC fraction is out of specification, but has a recovery of > 10%.

If any one surrogate in a fraction shows < 10 % recovery.

| All criteria were met |
|-----------------------|
| Criteria were not met |
| and/or see belowN/A |

VII. A MATRIX SPIKE/MATRIX SPIKE DUPLICATE (MS/MSD)

This data is generated to determine long term precision and accuracy in the analytical method for various matrices. This data alone cannot be used to evaluate the precision and accuracy of individual samples. If any % R in the MS or MSD falls outside the designated range, the reviewer should determine if there are matrix effects, i.e. LCS data are within the QC limits but MS/MSD data are outside QC limit.

1. MS/MSD Recoveries and Precision Criteria

The laboratory should use one MS and a duplicate analysis of an unspiked field sample if target analytes are expected in the sample. If target analytes are not expected, MS/MSD should be analyzed.

| | PD of the compounds | | | | ************************************** | |
|------------|------------------------|------------|---------|---------------------|--|------|
| MS OR MSD | COMPOUND | % R | RPD | QC LIMITS | ACTION | |
| | _are_not_required_as | • | | • | ike_used_to_assess | ; |
| * QC limit | s are laboratory in-ho | ouse perfo | mance o | criteria, LL = lowe | r limit, UL = upper lir | nit. |

Actions:

| QUALITY | %R < LL | %R > UL |
|--------------------|---------|---------|
| Positive results | J | J |
| Nondetects results | R | Accept |

If QC limits are not available, use limits of 70 - 130 %.

MS/MSD criteria apply only to the unspiked sample, its dilutions, and the associated MS/MSD samples:

If the % R for the affected compounds were < LL (or 70 %), qualify positive results (J) and nondetects (UJ).

If the % R for the affected compounds were > UL (or 130 %), only qualify positive results (J).

If 25 % or more of all MS/MSD %R were < LL (or 70 %) or if two or more MS/MSD %Rs were < 10%, qualify all positive results (J) and reject nondetects (R).

A separate worksheet should be used for each MS/MSD pair.

| All criteria were met |
|-----------------------|
| Criteria were not met |
| and/or see belowN/A |

VII. B MATRIX SPIKE/MATRIX SPIKE DUPLICATE

MS/MSD - Unspiked Compounds

It should be noted that Method TO-15 does not specify a MS/MSD criteria for the unspiked compounds in the sample. A %RSD of < 50% has therefore been utilized as professional judgment.

If all target analytes were spiked in the MS/MSD, this review element is not applicable.

List the %RSD of the compounds which do not meet the criteria.

| Sample ID: | | | Matrix/Level/Unit: | | | | |
|---|---|-------------|--------------------|----------|---|--|--|
| COMPOUND | SAMPLE CONC. | MS CONC. | MSD CONC. | % RSD | ACTION | | |
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Actions:

^{*} If the % RSD > 50, qualify the positive result in the unspiked samples as estimated (J).

^{*} If the % RSD is not calculated (NC) due to nondetected value, use professional judgment to qualify the data.

| All criteria were metX |
|------------------------|
| Criteria were not met |
| and/or see below |

VIII. LABORATORY CONTROL SAMPLE (LCS) ANALYSIS

This data is generated to determine accuracy of the analytical method for various matrices.

1. LCS Recoveries Criteria

Where LCS spiked with the same analyte at the same concentrations as the MS/MSD? Yes or No. If no make note in data review memo.

List the %R of compounds which do not meet the criteria

| FC2 ID | COMPOUND | % K | QC LIMI |
|-------------------------------------|--------------------------------|---------------------|---------------|
|)_(Blank_spike)_ _control_limits | _analyzed_in_this_data_package | e,_recoveries_a | nd_RPD_within |
| | | | |
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| | | | |

- QC limits are laboratory in-house performance criteria, LL = lower limit, UL = upper limit.
- * If QC limits are not available, use limits of 70 130 %.

Actions:

| QUALITY | %R < LL | %R > UL |
|--------------------|---------|---------|
| Positive results | J | J |
| Nondetects results | R | Accept |

All analytes in the associated sample results are qualified for the following criteria.

If 25 % of the LCS recoveries were < LL (or 70 %), qualify all positive results (j) and reject nondetects (R).

If two or more LCS were below 10 %, qualify all positive results as (J) and reject nondetects (R).

2. Frequency Criteria:

Where LCS analyzed at the required frequency and for each matrix? <u>Yes</u> or No. If no, the data may be affected. Use professional judgment to determine the severity of the effect and qualify data accordingly. Discuss any actions below and list the samples affected.

| | | | | All criteria were met Criteria were not met and/or see belowN/A |
|-----|-------------|---------------------|---------|---|
| IX. | LABORATOR | DUPLICATE PRECISION | | |
| | Sample IDs: | _B6-2IA/B6-2IA_Dup | Matrix: | Air |

Field duplicates samples may be taken and analyzed as an indication of overall precision. These analyses measure both field and lab precision; therefore, the results may have more variability than laboratory duplicates which only laboratory performance. It is also expected that soil duplicate results will have a greater variance than water matrices due to difficulties associated with collecting identical field duplicate samples.

The project QAPP should be reviewed for project-specific information.

Suggested criteria: RPD ± 25% for air samples. If both samples and duplicate are <5 SQL, the RPD criteria is doubled.

| COMPOUND | SQL | SAMPLE CONC. | DUPLICATE CONC. | RPD | ACTION |
|----------|--------------|--------------|------------------|-------------|-----------|
| | | | | | |
| RPI |) within the | e method per | formance criteri | a for all a | analytes. |
| | | | | | |
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Actions:

Qualify as estimated positive results (J) and nondetects (UJ) for the compound that exceeded the above criteria. For organics, only the sample and duplicate will be qualified.

If an RPD cannot be calculated because one or both of the sample results is not detected, the following actions apply:

If one sample result is not detected and the other is greater than 5x the SQL qualify (J/UJ).

If one sample value is not detected and the other is greater than 5x the SQL and the SQLs for the sample and duplicate are significantly different, use professional judgment to determine if qualification is appropriate.

If one sample value is not detected and the other is less than 5x, use professional judgment to determine if qualification is appropriate.

If both sample and duplicate results are not detected, no action is needed.

| All criteria | were | met | Χ |
|--------------|------|-----|---|
|--------------|------|-----|---|

| Criteria were not met |
|-----------------------|
| and/or see below |

X. INTERNAL STANDARD PERFORMANCE

The assessment of the internal standard (IS) parameter is used to assist the data reviewer in determining the condition of the analytical instrumentation.

List the internal standard area of samples which do not meet the criteria.

- * Area of +40% or -40% of the IS area in the associated calibration standard.
- * Retention time (RT) within \pm 0.06 seconds of the IS area in the associated calibration standard.

| DATE | SAMPLE ID | IS OUT | IS AREA | ACCEPTABLE RANGE | ACTION |
|----------|-------------------------------------|---|-------------------|----------------------|--------------|
| | tandard_area_and_reration_standards | etention_times_ | within_laboratory | _control_limits_for_ | both_samples |
| | | | | | |
| | | *************************************** | | | |
| Actions: | | | | | |

1. IS actions should be applied to the compound quantitated with the out-of-control ISs

| QUALITY | IS AREA < -40% | IS AREA > + 40% |
|---------------------|----------------|-----------------|
| Positive results | J | J |
| Nondetected results | R | ACCEPT |

2. If a IS retention time varies more than 0.330 seconds, the chromatographic profile for that sample must be examined to determine if any false positive or negative exists. For shifts of a large magnitude, the reviewer may consider partial or total rejection of the data for the sample fraction.

| All criteria were metX |
|------------------------|
| Criteria were not met |
| and/or see below |

XII. SAMPLE QUANTITATION

The sample quantitation evaluation is to verify laboratory quantitation results. In the space below, please show a minimum of one sample calculation:

1503058H-02A

2-Propanol

RF = 2.86097

[] = (230086)(25.0)/(94885)(2.86097)

= 153.983 ppbv OK

| All criteria were metX |
|------------------------|
| Criteria were not met |
| and/or see below |

XII. QUANTITATION LIMITS

A. Dilution performed

| SAMPLE ID | DILUTION FACTOR | REASONS FOR DILUTION |
|------------|-----------------|------------------------------|
| B6-1IA | 10.4 | High level of target species |
| B6-2IA | 6.37 | High level of target species |
| B6-2IA Dup | 6.53 | High level of target species |
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| 7.3 | | |
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| Percent Solids |
|---------------------------------------|
| List samples which have ≤ 50 % solids |
| |
| |

Actions:

If the % solids of a soil sample is 10-50%, estimate positive results (J) and nondetects (UJ)

If the % solids of a soil sample is < 10%, estimate positive results (J) and reject nondetects (R) $\,$

| Danie REVIEW OF VOLATILE ORGANI The following guidelines for evaluating volatile organics were | created to delineate required validations sional judgment to make more informed |
|---|---|
| | created to delineate required validations sional judgment to make more informed |
| actions. This document will assist the reviewer in using profest decision and in better serving the needs of the data users. The subset of JSEPA data validation guidance documents in the following ord D-1946 method for measuring permanent gases and light hydroxymples using gas chromatography (GC) and a thermal conductive detection (FID). Validating Air Samples. Volatile Organic Analyst (CO-15, (SOP # HW-31. Revision #4. October, 2006). The QC could be data review worksheets are from the primary guidance documents the hardcopied (laboratory name) _Eurofins | drocarbons in refinery and other source vity detector (TCD) and/or flame ionizations of Ambient Air in Canisters by Methoriteria and data validation actions listed of the ent, unless otherwise noted. data package received has been |
| _ab. Project/SDG No.:1503058F | Sample matrix:Air |
| No. of Samples:7 | Cample MauxAll |
| | |
| X Holding Times | _X Field Duplicates |
| N/A_GC/MS Tuning | _X Calibrations |
| | _X Compound Identifications |
| | _X Compound Quantitation |
| —————————————————————————————————————— | _X Quantitation Limits |
| N/A_Matrix Spike/Matrix Spike Duplicate | |
| Overall Comments:_Methane_by_ASTM_method_D-1946_(| (modified) |
| Definition of Qualifiers: J- Estimated results J- Compound not detected R- Rejected data JJ- Estimated nondetect | |
| Reviewer: | |

DATA COMPLETENESS

| MISSING INFORMATION | DATE LAB. CONTACTED | DATE RECEIVED |
|---|---------------------|---------------|
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| All criteria were metX |
|------------------------|
| Criteria were not met |
| and/or see below |

HOLDING TIMES

The objective of this parameter is to ascertain the validity of the results based on the holding time of the sample from time of collection to the time of analysis.

Complete table for all samples and note the analysis and/or preservation not within criteria

| SAMPLE ID | DATE SAMPLED | DATE ANALYZED | pН | ACTION |
|-----------|----------------------------|------------------------|---------|--------------|
| | | | | |
| | All a comba a comba a divi | :4h: - 4h | 1 | L -1.2 |
| <i></i> | All samples analyzed w | vitnin the recommended | metnoa_ | nolaing time |
| | | | | |
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Criteria

Aqueous samples – 14 days from sample collection for preserved samples (pH \leq 2, 4°C), no air bubbles.

Aqueous samples -7 days from sample collection for unpreserved samples, 4°C , no air bubbles.

Soil samples- 7 days from sample collection.

Cooler temperature (Criteria: 4 ± 2 °C): N/A – summa canisters

Actions

If the VOCs vial(s) have air bubbles, estimate positive results (J) and reject nondetects (R).

If the % solids of soil samples is 10-50%, estimates positive results (J) and nondetects (UJ)

If the % solid of soil samples is < 10%, estimate positive results (J) and reject nondetects (R).

If holding times are exceeded but < 14 days beyond criteria, estimate positive results (J) and nondetects (UJ).

If holding times are exceeded but < 28 days beyond criteria, estimate positive results (J) and reject nondetects (R).

If holding times are grossly exceeded (> 28 days beyond criteria), reject all results (R).

If samples were not iced or if the ice were melted (> 10°C), estimate positive results (J) and nondetects (UJ).

| | | | Il criteria were metN/A ere not met see below |
|--|----------------------------|-------------------------------------|--|
| GC/MS TUNING | | | |
| The assessment standard tuning (| | determine if the sample instrumen | tation is within the |
| N/A_ The BFB | performance results were | reviewed and found to be within the | specified criteria. |
| N/A_ BFB tuni | ng was performed for every | 24 hours of sample analysis. | |
| If no, use profes qualified or reject | | ine whether the associated data sl | nould be accepted, |
| List | the | samples | affected: |

If mass calibration is in error, all associated data are rejected.

Note: Samples analyzed using GC with either TCD or FID detection.

| All criteria were met _X |
|--------------------------|
| Criteria were not met |
| and/or see below |

CALIBRATION VERIFICATION

Compliance requirements for satisfactory instrument calibration are established to ensure that the instrument is capable of producing and maintaining acceptable quantitative data.

| Date of initial calibration: | 05/08/14 |
|------------------------------|----------------|
| Dates of continuing calibra | tion:_03/05/15 |
| Instrument ID numbers: | GC-9 |
| Matrix/Level: | Air/low |

| DATE | LAB ID# | FILE | CRITERIA OUT RFs, %RSD, %D, r | COMPOUND | SAMPLES AFFECTED |
|------|------------|---------------------------------------|-------------------------------------|-----------------------------|--------------------------|
| | | | rations meet method sprequirements. | pecific requirements. Initi | al calibration retention |
| | | · · · · · · · · · · · · · · · · · · · | | | |
| | | | | | |

Criteria

All RFs must be > 0.05 regardless of method requirements for SPCC.

All %RSD must be < 15 % regardless of method requirements for CCC.

All %Ds must be ≤ 30% regardless of method requirements for CCC.

Method TO-15 does not specify criterion for the curve correlation coefficient (r). A limit for r of \geq 0.995 has therefore been utilized as professional judgment.

Actions

If any compound has an initial RF or a continuing RF of < 0.05, estimate positive results (J) and reject nondetects (R), regardless of method requirements.

If any compound has a %RSD > 15%, estimate positive results (J) and use professional judgment to qualify nondetects.

If any compound has a %RSD > 90%, estimate positive results (J) and reject nondetects (R).

If any compound has a % D > 30%, estimate positive results (J) and reject nondetects (R).

If any compound has a % D > 30%, estimate positive results (J) and nondetects (UJ).

If any compound has a % D > 90%, estimate positive results (J) and reject nondetects (R).

If any compound has r < 0.995, estimate positive results and nondetects.

A separate worksheet should be filled for each initial curve

| All criteria were metX |
|------------------------|
| Criteria were not met |
| and/or see below |

V A. BLANK ANALYSIS RESULTS (Sections 1 & 2)

The assessment of the blank analysis results is to determine the existence and magnitude of contamination problems. The criteria for evaluation of blanks apply only to blanks associated with the samples, including trip, equipment, and laboratory blanks. If problems with any blanks exist, all data associated with the case must be carefully evaluated to determine whether or not there is an inherent variability in the data for the case, or if the problem is an isolated occurrence not affecting other data.

List the contamination in the blanks below. High and low levels blanks must be treated separately.

Laboratory blanks

| DATE ANALYZED | LAB ID | LEVEL/ MATRIX | COMPOUND | CONCENTRATION UNITS |
|------------------|-----------------|-------------------|---------------------|---------------------|
| All_method | d_blank_meeth_ | _method_speci | fic_criteria | |
| Summa_c | anisters_met_cl | eaning_certifica | ation_criteria | |
| Field/Equipmen | t/Trip blank | | | |
| DATE ANALYZED | LAB ID | LEVEL/ MATRIX | COMPOUND | CONCENTRATION UNITS |
| No_field/trip/eq | uipment_blanks | _analyzed_with | n_this_data_package | |
| | | | | |
| | | | | |
| | | | | |
| | | | | |

| All criteria were metX |
|------------------------|
| Criteria were not met |
| and/or see below |

VB. BLANK ANALYSIS RESULTS (Section 3)

Blank Actions

Action Levels (ALs) should be based upon the highest concentration of contaminant determined in any blank. Do not qualify any blank with another blank. The ALs for samples which have been diluted should be corrected for the sample dilution factor and/or % moisture, where applicable. No positive sample results should be reported unless the concentration of the compound in the samples exceeds the ALs:

ALs = 10x the amount of common contaminants (methylene chloride, acetone, 2-butanone, and toluene)

ALs = 5x for any other compounds

Specific actions are as follows:

If the concentration is < sample quantitation limit (SQL) and \le AL, report the compound as not detected (U) at the SQL.

If the concentration is \geq SQL but \leq AL, report the compound as not detected (U) at the reported concentration.

If the concentration is \geq SQL and > AL, report the concentration unqualified.

Notes:

High and low level blanks must be treated separately Compounds qualified "U" for blank contamination are still considered "hits" when qualifying for calibration criteria.

| CONTAMINATION SOURCE/LEVEL | COMPOUND | CONC/UNITS | AL/UNITS | SQL | AFFECTED SAMPLES |
|----------------------------|----------|------------|----------|-----|------------------|
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| All criteria were met _ | _N/A |
|-------------------------|------|
| Criteria were not met | |
| and/or see below | |

ACTION

SURROGATE SPIKE RECOVERIES

Laboratory performance of individual samples is established by evaluation of surrogate spike recoveries. All samples are spiked with surrogate compounds prior to sample analysis. The accuracy of the analysis is measured by the surrogate percent recovery. Since the effects of the sample matrix are frequently outside the control of the laboratory and may present relatively unique problems, the validation of data is frequently subjective and demands analytical experience and professional judgment.

SURROGATE COMPOUND

List the percent recoveries (%Rs) which do not meet the criteria for surrogate recovery. Matrix: solid/aqueous

| _Surrogate_standard | ds_not_requ | iired_by_the_me | thod | | |
|------------------------------|-------------|---|------|----|--|
| | | | | | |
| | | *************************************** | | | |
| QC Limits* (Air) LL_to_UL | to | to | to | to | |

- * QC limits are laboratory in-house performance criteria, LL = lower limit, UL = upper limit.
- * If QC limits are not available, use limits of 80 120 % for aqueous and 70 130 % for solid samples.

Actions:

SAMPLE ID

| QUALITY | %R < 10% | %R = 10% - LL | %R > UL |
|--------------------|----------|---------------|---------|
| Positive results | J | J | J |
| Nondetects results | R | UJ | Accept |

Surrogate action should be applied:

If one or more surrogate in the VOC fraction is out of specification, but has a recovery of > 10%.

If any one surrogate in a fraction shows < 10 % recovery.

| All criteria were met |
|-----------------------|
| Criteria were not met |
| and/or see belowN/A |

VII. A MATRIX SPIKE/MATRIX SPIKE DUPLICATE (MS/MSD)

This data is generated to determine long term precision and accuracy in the analytical method for various matrices. This data alone cannot be used to evaluate the precision and accuracy of individual samples. If any % R in the MS or MSD falls outside the designated range, the reviewer should determine if there are matrix effects, i.e. LCS data are within the QC limits but MS/MSD data are outside QC limit.

1. MS/MSD Recoveries and Precision Criteria

The laboratory should use one MS and a duplicate analysis of an unspiked field sample if target analytes are expected in the sample. If target analytes are not expected, MS/MSD should be analyzed.

List the %Rs, RPD of the compounds which do not meet the criteria.

| Sample ID: | | | Matrix/Level: | | |
|------------|-----------------------|-----|---------------|-----------------|------------------------|
| MS OR MSD | COMPOUND | % R | RPD | QC LIMITS | ACTION |
| | _not_required_as_part | | 1-method | I_D-1946;_blanl | <_spike_used_to_assess |

* If QC limits are not available, use limits of 70 – 130 %.

Actions:

| QUALITY | %R < LL | %R > UL |
|--------------------|---------|---------|
| Positive results | J | J |
| Nondetects results | R | Accept |

MS/MSD criteria apply only to the unspiked sample, its dilutions, and the associated MS/MSD samples:

If the % R for the affected compounds were < LL (or 70 %), qualify positive results (J) and nondetects (UJ).

If the % R for the affected compounds were > UL (or 130 %), only qualify positive results (J).

If 25 % or more of all MS/MSD %R were < LL (or 70 %) or if two or more MS/MSD %Rs were < 10%, qualify all positive results (J) and reject nondetects (R).

A separate worksheet should be used for each MS/MSD pair.

^{*} QC limits are laboratory in-house performance criteria, LL = lower limit, UL = upper limit.

| All criteria were met |
|-----------------------|
| Criteria were not met |
| and/or see below N/A |

VII. B MATRIX SPIKE/MATRIX SPIKE DUPLICATE

MS/MSD - Unspiked Compounds

It should be noted that Method TO-15 does not specify a MS/MSD criteria for the unspiked compounds in the sample. A %RSD of < 50% has therefore been utilized as professional judgment.

If all target analytes were spiked in the MS/MSD, this review element is not applicable.

List the %RSD of the compounds which do not meet the criteria.

| Sample ID: | | | Matrix/Level/Unit: | | | |
|------------|-----------------|----------|--------------------|-------|--------|--|
| COMPOUND | SAMPLE CONC. | MS CONC. | MSD CONC. | % RSD | ACTION | |
| | | | | | | |
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Actions:

^{*} If the % RSD > 50, qualify the positive result in the unspiked samples as estimated (J).

^{*} If the % RSD is not calculated (NC) due to nondetected value, use professional judgment to qualify the data.

| All criteria were metX |
|------------------------|
| Criteria were not met |
| and/or see below |

VIII. LABORATORY CONTROL SAMPLE (LCS) ANALYSIS

This data is generated to determine accuracy of the analytical method for various matrices.

1. LCS Recoveries Criteria

Where LCS spiked with the same analyte at the same concentrations as the MS/MSD? Yes or No. If no make note in data review memo.

List the %R of compounds which do not meet the criteria

| | LCS ID | COMPOUND | % R | QC LIMIT |
|---------|-------------------|---------------------------|---------------------------------------|----------|
| LCS/L | CSD_(Blank_spik | ce)_analyzed_in_this_data | _package;_recoveries_ | and_RPD |
| within_ | _laboratory_contr | ol_limits | | |
| | | | | |
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| | | | | |

- * QC limits are laboratory in-house performance criteria, LL = lower limit, UL = upper limit.
- * If QC limits are not available, use limits of 70 130 %.

Actions:

| QUALITY | %R < LL | %R > UL |
|--------------------|---------|---------|
| Positive results | J | J |
| Nondetects results | R | Accept |

All analytes in the associated sample results are qualified for the following criteria.

If 25 % of the LCS recoveries were < LL (or 70 %), qualify all positive results (j) and reject nondetects (R).

If two or more LCS were below 10 %, qualify all positive results as (J) and reject nondetects (R).

2. Frequency Criteria:

Where LCS analyzed at the required frequency and for each matrix? <u>Yes</u> or No. If no, the data may be affected. Use professional judgment to determine the severity of the effect and qualify data accordingly. Discuss any actions below and list the samples affected.

| | | All criteria were metX Criteria were not met and/or see below |
|-----|--|---|
| IX. | FIELD/LABORATORY DUPLICATE PRECISION | |
| | Sample ID_B6-2IA/B6-2IA_Dup Sample ID_B6-2SS/B6-2SS_Dup | Matrix:Air Matrix:Air |

Field/laboratory duplicates samples may be taken and analyzed as an indication of overall precision. These analyses measure both field and lab precision; therefore, the results may have more variability than laboratory duplicates which only laboratory performance. It is also expected that soil duplicate results will have a greater variance than water matrices due to difficulties associated with collecting identical field duplicate samples.

The project QAPP should be reviewed for project-specific information. Suggested criteria: RPD \pm 25% for air samples. If both samples and duplicate are <5 SQL, the RPD criteria is doubled.

| COMPOUND | SQL | SAMPLE CONC. | DUPLICATE CONC. | RPD | ACTION |
|--------------------------|-----------|--------------|-------------------------------------|-----|-----------------------------|
| RPD for field duplicates | within la | | trol limits. RPD tatory control lim | | ratory duplicate (LCS/LCSD) |
| | | | | | |

Actions:

Qualify as estimated positive results (J) and nondetects (UJ) for the compound that exceeded the above criteria. For organics, only the sample and duplicate will be qualified.

If an RPD cannot be calculated because one or both of the sample results is not detected, the following actions apply:

If one sample result is not detected and the other is greater than 5x the SQL qualify (J/UJ).

If one sample value is not detected and the other is greater than 5x the SQL and the SQLs for the sample and duplicate are significantly different, use professional judgment to determine if qualification is appropriate.

If one sample value is not detected and the other is less than 5x, use professional judgment to determine if qualification is appropriate.

If both sample and duplicate results are not detected, no action is needed.

| All criteria were metN/A |
|--------------------------|
| Criteria were not met |
| and/or see below |

X. INTERNAL STANDARD PERFORMANCE

The assessment of the internal standard (IS) parameter is used to assist the data reviewer in determining the condition of the analytical instrumentation.

List the internal standard area of samples which do not meet the criteria.

- * Area of +40% or -40% of the IS area in the associated calibration standard.
- * Retention time (RT) within \pm 0.06 seconds of the IS area in the associated calibration standard.

| DATE | SAMPLE ID | IS OUT | IS AREA | ACCEPTABLE RANGE | ACTION |
|-------------|----------------------|-------------|---------|---------------------------------------|---|
| | tandard_not_required | - | | ntified_by_externa | l_standard |
| ··· | | | | | |
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| Actions: | | | | | |

1. IS actions should be applied to the compound quantitated with the out-of-control ISs

| QUALITY | IS AREA < -40% | IS AREA > + 40% |
|---------------------|----------------|-----------------|
| Positive results | J | J |
| Nondetected results | R | ACCEPT |

2. If a IS retention time varies more than 0.330 seconds, the chromatographic profile for that sample must be examined to determine if any false positive or negative exists. For shifts of a large magnitude, the reviewer may consider partial or total rejection of the data for the sample fraction.

| All criteria were metX |
|------------------------|
| Criteria were not met |
| and/or see below |

XII. SAMPLE QUANTITATION

The sample quantitation evaluation is to verify laboratory quantitation results. In the space below, please show a minimum of one sample calculation:

1503058F-01A

Methane

RF = 159808871

[] = (17549)/(159808871)

= 0.00011 % OK

| All criteria were metX |
|------------------------|
| Criteria were not met |
| and/or see below |

| XII. | ΩL | IANTI | TAT | ION. | LIMITS | 3 |
|-------|----|-------------|------------|------|--------|---|
| /\II. | w | // (1 4 1 1 | $I \cap I$ | | | _ |

A. Dilution performed

| SAMPLE ID | DILUTION FACTOR | REASONS FOR DILUTION | | | | | |
|------------------|-----------------------------|----------------------|--|--|--|--|--|
| All samples dilu | ited by a factor of less th | | | | | | |
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| Pe | rcent S | olids | | | | |
|-----|---------|-----------|---------------------|----------|------|---|
| Lis | t samp | les which | have <u><</u> 50 | % solids | | |
| | · | | _ | | | |
| | | | | | | |
| | | | | | | · |

Actions:

If the % solids of a soil sample is 10-50%, estimate positive results (J) and nondetects (UJ)

If the % solids of a soil sample is < 10%, estimate positive results (J) and reject nondetects (R)